

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

⦿ **BLACK BORDERS**

- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.**

PATENT SPECIFICATION

NO DRAWINGS

828,880



Date of Application and filing Complete Specification July 7, 1958.

No. 21721/58.

Application made in United States of America on July 15, 1957.

Complete Specification Published Feb. 24, 1960.

Index at acceptance: —Classes 2(3), C2A(3:5:14), C2R15; and 81(1), B2(B3:G:L:N:P:S).

International Classification: —A61k, C07c.

COMPLETE SPECIFICATION

**2-Amino-1-(3,4-Methylenedioxyphenyl)-Propane Isomers and an
Ataractic preparation containing 2-Amino-1-(3,4-
Methylenedioxyphenyl)-Propane**

We, SMITH KLINE & FRENCH LABORATORIES, a Corporation organized under the Laws of the State of Delaware, one of the United States of America, of 1530, Spring Garden Street, City of Philadelphia, Pennsylvania, United States of America, do hereby declare the invention, for which we pray that a patent may be granted to us, and the method by which it is to be performed, to be particularly described in and by the following statement:—

This invention relates to novel isomers of 2-amino-1-(3,4-methylenedioxyphenyl)-propane, and to a medicinal preparation having ataractic activity.

Prior to the present invention the important advances in the treatment of mentally deranged have largely been in the excited group of patients through the use of central nervous system depressant compounds commonly referred to as tranquilizers. A large proportion of the population of mental hospitals, however, consists of depressed patients whose conditions generally are either not responsive to tranquilizers or aggravated by the use of these drugs. The need of a safe, effective composition for use in this area has been great.

The preparation in accordance with this invention contains 2-amino-1-(3,4-methylenedioxyphenyl)-propane and is very useful in treating various depressive states of psychotic patients due to having an unusual differential in its activity. It, surprisingly for a central nervous stimulant, provides a strong conditioned response block in animals. In the treatment of severely depressed psychotics, it induces ataraxia without any substantial amount of the sympathomimetic action found in closely related compounds such as amphetamine. This preparation has a low incidence of side effects in a dosage range where preparations containing closely related

compounds such as 2-amino-1-phenylpropanes produce severe side effects such as jitteriness, excessive stimulation or increased tension.

More specifically, the preparation of this invention is in a dosage unit form and comprises from about 15 mg. to about 150 mg., and preferably from about 25 mg. to about 100 mg., of 2-amino-1-(3,4-methylenedioxyphenyl)-propane or a non-toxic acid solution salt thereof and a pharmaceutical carrier.

The *d*- or *l*-isomer of 2-amino-1-(3,4-methylenedioxyphenyl)-propane or a non-toxic salt thereof can be substituted advantageously for the racemic mixture. Where the term 2-amino-1-(3,4-methylenedioxyphenyl)-propane is employed without any indication as to the *d*-, *l*- or racemic form, it is intended herein and in the claims to cover the individual *d*- and *l*-isomers as well as mixtures thereof.

The *l*-isomer is advantageous since it also is an effective anorexic agent and, hence, its employment is advantageous where it is desired to curb the appetite.

The active *d*-isomer is prepared by dissolving the racemic hydrochloride salt in water, neutralizing with an inorganic base, for example, sodium hydroxide, and extracting into an organic solvent such as ether or benzene. *d*-Tartaric acid is added to separate the *d*-tartrate salt. Recrystallization from alcohol such as isopropanol or aqueous isopropanol gives the pure *d*-isomer as the *d*-tartrate with an optical rotation of 29.4° (2% in water). The *d*-base in hexane has a rotation of 24.6° (1%). If desired, the hydrochloride salt may be regenerated from the active base by treating an ether or hexane solution with anhydrous hydrogen chloride gas. The *l*-base is similarly prepared.

Preferably the hydrochloric salt of the 2-amino-1-(3,4-methylenedioxyphenyl)-

[Pr]

propane is used, however, either the base itself or a non-toxic pharmaceutically acceptable acid addition salt of the base may be used, such as the salt derived from sulfuric, nitric, phosphoric, citric, acetic, lactic, salicylic, tartaric, ethanedisulfonic, sulfamic, acetylsalicylic, succinic, fumaric, maleic, hydrobromic, or benzoic acid. The salts are conveniently prepared by reacting the free base with either a stoichiometric amount or an excess of the desired acid in a suitable solvent such as ethanol, ether, ethyl acetate, acetone, water or various combinations of solvents.

The lower part of the dosage range of the 2-amino-1-(3,4-methylenedioxyphenyl)-propane of from about 15 mg. to about 25 mg. is aimed at child medication and at parenteral preparations. For oral use with a solid carrier the preparation for adults would advantageously contain from about 25 mg. to about 75 mg. of the active propane compound. If a sustained release (i.e. having a release over a period of about 12 hours) is used, the above dosage ranges can be tripled.

The pharmaceutical carrier may be, for example, either a solid or a liquid. Exemplary of solid carriers are talc, corn starch, lactose, ethylcellulose, magnesium stearate, agar, pectin, stearic acid, gelatin and acacia. Exemplary of liquid carriers are water, peanut oil, olive oil and sesame oil. Solid carriers are preferred.

A wide variety of pharmaceutical forms can be employed. Thus, if a solid carrier is used, the preparation can be tableted or placed in a hard gelatin capsule. If a liquid carrier is used, the preparation may be in the form of a soft gelatin capsule or placed in an ampule. The amount of carrier will vary widely but preferably will be from about 25 mg. to about 1 gm.

The preparation of this invention may be administered internally in an amount to produce ataraxia in depressed psychotic patients. The administration may be orally or parenterally preferably employing the above described preparation. In this method it is preferred to administer from about 60 mg. to about 350 mg. and advantageously about 75 mg. to about 320 mg. of 2-amino-1-(3,4-methylenedioxyphenyl)-propane or a salt thereof daily, preferably administering equal doses three or four times daily. In the treatment of children somewhat lower dosages are used depending largely on the age and weight of the child. Such doses may be individually determined by the physician but will ordinarily be about half the adult dosage.

60 EXAMPLE I

A solution of 35.8 g. (0.2 mole) of 2-amino-1-(3,4-methylenedioxyphenyl)-propane and 30 g. of *d*-tartaric acid in 600 ml. of 75% isopropanol is allowed to stand at room tempera-

ture after seeding. A thick precipitate separates. After filtration, the solid tartrate is recrystallized several times from isopropanol to white crystals of *d*-2-amino-1-(3,4-methylenedioxyphenyl)-propane *d*-tartrate, m.p. 145–146° C., $[\alpha]^{25}_D$ and 29.44° (1% H₂O). The free *d*-base is regenerated and taken into hexane, $[\alpha]^{25}_D$ +24.6°. The free *d*-base is reconverted to the hydrochloride salt with gaseous hydrogen chloride, m.p. 185–187° C.

The mother filtrate is evaporated to give 22 g. of the 1-2-amino-1-(3,4-methylenedioxyphenyl)-propane *d*-tartrate, m.p. 125–130° C. After converting a portion to the base in hexane, the specific rotation of this sample is –11.5° C. The remainder of the tartrate is recrystallized from aqueous ethanol to pure white crystals of *l*-base *d*-tartrate, m.p. 129–132° C., $[\alpha]^{25}_D$ –28.5° (1% H₂O).

EXAMPLE 2

dl-2-Amino-1-(3,4-methylene-dioxyphenyl)-propane
Hydrochloride - 25 mg.
Lactose - 230 mg.
Starch - 45 mg.

The above ingredients were thoroughly mixed, granulated using a 10% gelatin solution and compressed into tablets using an admixture of talc-stearic acid as a lubricant.

EXAMPLE 3

dl-2-Amino-1-(3,4-methylene-dioxyphenyl)-propane
Maleate - 75 mg.
Lactose - 225 mg.

The above ingredients were thoroughly mixed, granulated using a 50% sucrose solution and compressed into tablets using an admixture of 7% starch and 1% magnesium stearate based on tablet weight.

EXAMPLE 4

d-2-Amino-1-(3,4-methylene-dioxyphenyl)-propane
Hydrochloride - 50 mg.
Lactose - 150 mg.
Starch - 50 mg.

The above ingredients were thoroughly mixed, granulated using a 10% gelatin solution and compressed into scored tablets.

EXAMPLE 5

dl-2-Amino-1-(3,4-methylene-dioxyphenyl)-propane
Hydrochloride - 300.00 gm.
Lactose
(200 mesh) - 2820.00 gm.
Magnesium stearate - 60.00 gm.

The powders are mixed, screened and filled into No. 2 hard gelatin capsules (12,000 capsules at 25 mg).

EXAMPLE 6

- 1 - 2 - Amino 1 - (3,4 - methylene -
dioxiphenyl)-propane
Sulfate - - - 75 mg.
5 Peanut oil - - - 225 mg.
The ingredients are mixed to a thick slurry
and filled into a soft gelatin capsule.

EXAMPLE 7

- 10 dl - 2 - Amino - 1 - (3,4 - methylene -
dioxiphenyl)-propane
Hydrochloride - - 100 mg.
Hydrogenated castor
oil - - - 100 mg.

15 The chemical is imbedded in the hydro-
genated castor oil by melting the latter, mix-
ing in the chemical and solidifying. After
comminuting and screening through a Num-
ber 10 screen, the powder is granulated with
a small amount of starch to produce sustained
20 release granules.

- dl - 2 - Amino - 1 - (3,4 - methylene -
dioxiphenyl)-propane
Hydrochloride - - 50 mg.
25 Stearic acid - - - 15 mg.
Talc - - - 15 mg.

The above ingredients are mixed and
granulated with a gelatin solution, dried,
30 screened and compressed into cylindrical,
flat faced tablets. The sustained release
granules are added to the die and compressed
onto the previously formed tablets.

EXAMPLE 8

- d - 2 - Amino - 1 - (3,4 - methylene -
dioxiphenyl)-propane
35 Hydrochloride - - 15 mg.
Lactose - - - 245 mg.
Magnesium stearate 5 mg.

The powders are mixed, screened and filled
into a Number 2 hard gelatin capsule.

EXAMPLE 9

- 40 dl - 2 - Amino - 1 - (3,4 - methylene -
dioxiphenyl)-propane
Hydrochloride - - 30 mg.
Lactose - - - 225 mg.
45 Starch - - - 45 mg.

The ingredients are mixed, granulated and
compressed into a scored tablet which may
be broken for divided doses if desired.

EXAMPLE 10

- dl - 2 - Amino - 1 - (3,4 - methylene - 50
dioxiphenyl)-propane
Hydrochloride - - 2.0 w/v
Sodium chloride - 0.375 w/v
Water for injection,
U.S.P., q.s. ad 100 % 55

The solid ingredients are dissolved in part
of the water and made to 100% volume. The
resulting solution is filtered through a Selas
filter and filled into ampuls. The word
"Selas" is a registered Trade Mark. 60

WHAT WE CLAIM IS:—

1. A pharmaceutical preparation having
ataractic activity, in dosage unit form, com-
prising a pharmaceutical carrier and a 2-
amino 1 - (3,4 - methylenedioxiphenyl) - 65
propane or its non-toxic acid addition salts.
2. The preparation claimed in Claim 1 in
which the dosage unit form is a capsule.
3. The preparation claimed in Claim 1 in
which the dosage unit form is a tablet. 70
4. The preparation claimed in any of Claims
1 to 3 in which the 2-amino-1-(3,4-methylene-
dioxiphenyl)-propane is in the racemic form.
5. The preparation claimed in any of Claims
1 to 3 in which the 2-amino-1-(3,4-methylene- 75
dioxiphenyl)-propane is in the dextro isomer.
6. The preparation claimed in any of
Claims 1 to 3 in which the 2-amino-1-(3,4-
methylenedioxiphenyl)-propane is the levo
isomer. 80
7. The preparation claimed in any of the
preceding claims in which the 2-amino-1-(3,4-
methylenedioxiphenyl)-propane or its non-
toxic acid addition salts are present in an
amount of from about 15 mg to about 150 mg. 85
8. The preparation claimed in any of
Claims 1 to 6 in which the 2-amino-1-(3,4-
methylenedioxiphenyl)-propane or its non-
toxic acid addition salts are present in an
amount of from about 25 mg. to about 100 90
mg.
9. d - 2 - Amino - 1 - (3,4 - methylene -
dioxiphenyl) - propane or its non-toxic acid
addition salts.
10. l - 2 - Amino - 1 - (3,4 - methylene - 95
dioxiphenyl)-propane or its non-toxic acid
addition salts.

HASELTINE, LAKE & CO.,
28, Southampton Buildings, London, W.C.2,
Agents for the Applicants.